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Published in:
Nature Communications

DOI:
[10.1038/ncomms9079](https://doi.org/10.1038/ncomms9079)

Publication date:
2015

Citation for published version (APA):

Bennett, T. D., Tan, J., Yue, Y., Baxter, E., Ducati, C., Terrill, N. J., Yeung, H. H. -M., Zhou, Z., Chen, W., Henke, S., Cheetham, A. K., & Greaves, G. N. (2015). Hybrid glasses from strong and fragile metal-organic framework liquids. *Nature Communications*, 6, 1-7. [8079]. <https://doi.org/10.1038/ncomms9079>

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Received 15 Jun 2015 | Accepted 15 Jul 2015 | Published 28 Aug 2015

DOI: 10.1038/ncomms9079

OPEN

Hybrid glasses from strong and fragile metal-organic framework liquids

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Hybrid glasses connect the emerging field of metal-organic frameworks (MOFs) with the glass formation, amorphization and melting processes of these chemically versatile systems. Though inorganic zeolites collapse around the glass transition and melt at higher temperatures, the relationship between amorphization and melting has so far not been investigated. Here we show how heating MOFs of zeolitic topology first results in a low density 'perfect' glass, similar to those formed in ice, silicon and disaccharides. This order-order transition leads to a super-strong liquid of low fragility that dynamically controls collapse, before a subsequent order-disorder transition, which creates a more fragile high-density liquid. After crystallization to a dense phase, which can be remelted, subsequent quenching results in a bulk glass, virtually identical to the high-density phase. We provide evidence that the wide-ranging melting temperatures of zeolitic MOFs are related to their network topologies and opens up the possibility of 'melt-casting' MOF glasses.

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The microporous hybrid materials known as metal-organic frameworks (MOFs) consist of inorganic clusters or ions bridged by organic ligands in open, as well as dense three-dimensional arrays. The former are of great interest owing to their potential use in gas separation and storage, and the latter in multiferroic, conductive and drug/harmful waste encapsulation applications^{1–3}. An important subset of MOFs, the zeolitic imidazolate frameworks (ZIFs), adopt similar network structures to zeolites (inorganic low-density frameworks of corner sharing SiO_4 and AlO_4 tetrahedra), and, in particular, undergo thermal and pressure-induced amorphization (loss of periodicity)^{4–7}. High-density amorphous (HDA) inorganic glasses, along with low-density amorphous (LDA) states, of identical topologies to their parent crystalline phases have previously been identified via zeolite amorphization^{4,6}.

Such LDA states are also referred to as ‘perfect’ glasses⁸, and were first observed by depressurizing pressure-induced HDA phases or desolvating crystalline structures in, for example, ice⁹, silicon¹⁰ and trehalose¹¹. LDA phases are of scientific interest because of their location deep in the potential energy landscape (PEL)^{12,13}, at similar potential energies to their crystalline equivalents. This is in contrast to HDA phases that share the same composition as their perfect glass LDA counterparts, have greater entropy as well as density, and are located higher in the PEL. Moreover, compared with HDA phases, LDA phases have unique mechanical properties^{12,14}, which are connected to the formation of ultrastable glasses¹⁵. In particular, perfect glasses have been predicted to soften to super-strong liquids (low-density liquid (LDLs))⁸ above the glass transition temperature T_g in the supercooled state.

Experimentally, for most glass systems $T_g \sim 2/3 T_m$, defining the practical limits of the supercooled state¹². For glasses with a well-defined T_g , but that happen to decompose on heating before they melt, this relationship offers the opportunity to project a ‘virtual melting point’ that can be compared with the actual melting points T_m of isomorphous systems that survive the transition at T_m from the supercooled to the liquid state.

The dynamic behaviour of a supercooled liquid is quantified through the fragility index, m , equation (1), which measures, on a reduced temperature scale, the activation energy of the viscosity η at the glass transition T_g (ref. 16). T_g is defined to occur when η reaches 10^{12} Pa.s. While silica is the strongest liquid among conventional glass-forming liquids, fragilities for some LDL phases fall between 12 (ref. 13) and 14 (ref. 17), endorsing them as super-strong liquids, the antecedents of perfect glasses^{7,12}.

$$m = \left[\frac{d(\log \eta)}{d\left(\frac{T_g}{T}\right)} \right]_{T=T_g} \quad (1)$$

In the context of forming glasses from the collapse of zeolitic structures, the prospect was that this might lead to glasses sharing similar topology to precursor crystals¹². Indeed the ordered nature of the LDA perfect (low entropy) glass phase was identified by the retention of the THz features defining zeolitic topology when the majority of the starting crystal had amorphized⁶. This was in contrast to the eventual emergence of a featureless boson peak at higher temperatures, typical of less well-ordered conventional higher entropy HDA glasses¹². Glass transition temperatures of LDA phases were found to be significantly greater than HDA phases with fragilities in the super-strong range^{4,13}. Finally, by combining temperature- and pressure (P)-induced amorphization experiments, a critical point was identified at negative pressure¹³ and a negative dT/dP slope for the LDL–high-density liquid (HDL) transition. Considering

the Clapeyron relation, $dT/dP = \Delta V/\Delta S$, if dT/dP is negative, an increase in entropy (S) signifies a decrease in volume (V) and increase in density, endorsing the LDA and HDA assignments that had been made.

In the present work, we turn our attention to MOFs, research of the glassy behaviour of which is scarce^{18,19}. Specifically, we contrast ZIF-4 [$\text{Zn}(\text{C}_3\text{H}_3\text{N}_2)_2$] with ZIF-8 [$\text{Zn}(\text{C}_4\text{H}_5\text{N}_2)_2$] (ref. 20). We study the mechanism of amorphization of ZIF-4 by thermogravimetric analysis, differential scanning calorimetry (DSC), X-ray total scattering and *in situ* small- and wide-angle X-ray scattering (small angle X-ray scattering (SAXS)/wide angle X-ray scattering (WAXS)) experiments. Importantly, the different T_g s of the LDA, HDA and melt-quenched hybrid glass (MQG) reflect their differing depths in the PEL and the differences in fragility of the corresponding supercooled liquids. While ZIF-8 decomposes before it melts, the ‘virtual’ T_m discussed above can be calculated, lying close to the ‘real’ T_m of its inorganic counterpart. This suggests the dominance of network architecture in melting, characterized by collective THz vibrations⁶,

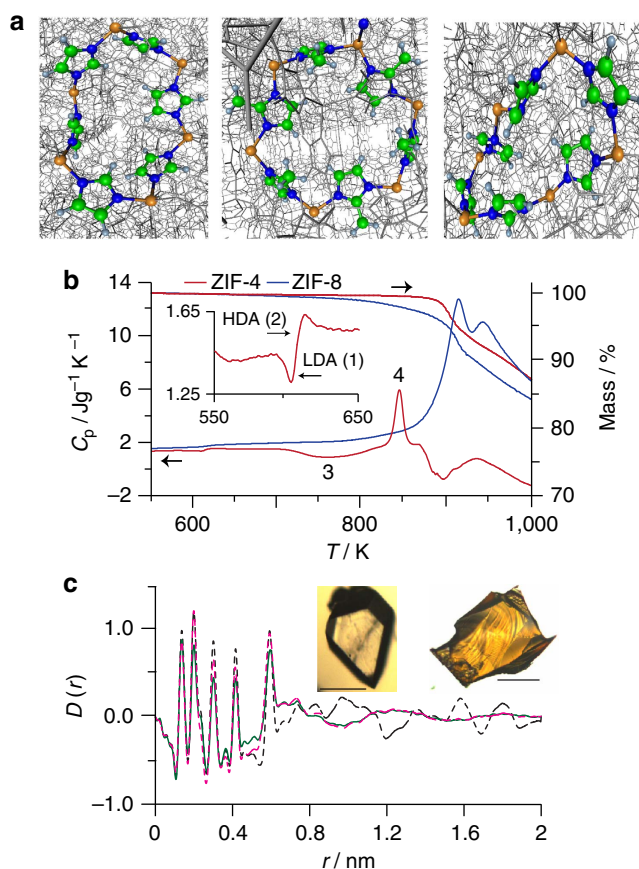


Figure 1 | Phase transitions of ZIF-4 on heating. (a) Highlighting the rings and imidazolate linkages in zeolitic topologies, in the ordered structure of crystalline ZIF-4 (left) and ZIF-8 (center), and the disordered HDA phase (right) obtained by Molecular Dynamics modelling (Supplementary Methods). Zn, orange; N, blue; C, green; and H, grey. (b) Thermogravimetric analysis and C_p plots for ZIF-4 and ZIF-8, showing, for the former (inset), exothermic collapse to the LDA phase (1) which is closely followed by (2) endothermic formation of the HDA phase, and (3) recrystallization (exothermic). Endothermic melting (4) then follows before thermal degradation. (c) X-ray PDF data $D(r)$ measured for the MQG (green), ZIF-4 (broken black) and the HDA phase (broken pink). The X-ray total scattering data $S(q)$ is presented in Supplementary Fig. 1. Inset: optical images of (left) ZIF-4 (right) MQG, showing the typical fracture pattern of a non-metallic bulk glass. Scale bars, 100 μm .

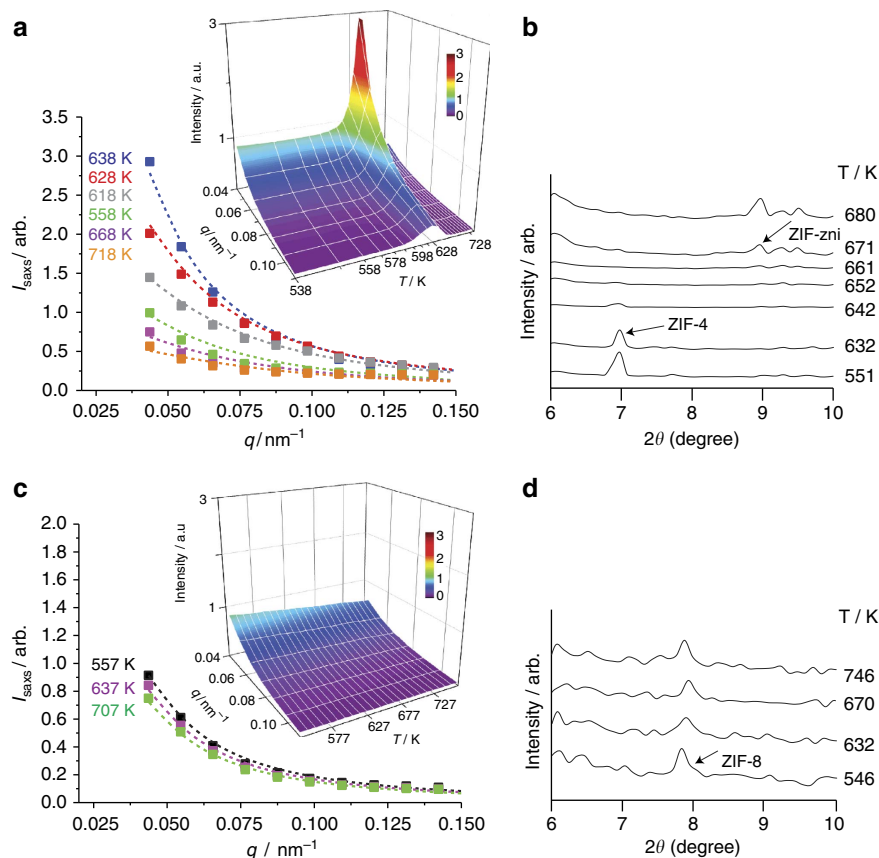


Figure 2 | SAXS/WAXS data on ZIF-4 (top) and ZIF-8 (bottom). (a) $I(q)_{\text{SAXS}}$ profiles of ZIF-4, with Lorentzian fits (Supplementary Methods) and three-dimensional plot (inset), highlighting the emergence of a peak between 618 and 663 K (Supplementary Fig. 2f). (b) WAXS data shows the major loss of Bragg diffraction on collapse at ca. 642 K. (c) $I(q)_{\text{SAXS}}$ profiles with Lorentzian fits and three-dimensional plot of the SAXS results for ZIF-8. (d) WAXS data show the retention of crystallinity across the entire temperature range studied.

in contrast to the simplistic interatomic variance condition for melting enshrined in Lindemann's law²¹.

Results

Differential scanning calorimetry. ZIF-4 collapses to an HDA phase, through formation of a LDA phase, before recrystallization into the dense ZIF-zni structure, which can be remelted. Quenching from below the decomposition temperature leads to the formation of a bulk MQG, virtually indistinguishable from the HDA phase, despite each having totally different thermal histories (Fig. 1). T_m for ZIF-4 lies close to that of an inorganic phosphate with a related zeolitic topology²². ZIF-8, on the other hand, adopts the sodalite structure, and does not thermally collapse, though amorphizes under pressure²³.

Small and wide angle X-ray scattering. Variable temperature SAXS and WAXS measurements were performed to probe the mechanism of amorphization. The SAXS signal $I(q)_{\text{SAXS}}$, which measures differences in local density¹², crucially continues after the majority of Bragg diffraction disappears, the SAXS maximum extending to significantly higher temperatures (Fig. 2a,b and Supplementary Fig. 2), as found earlier for conventional zeolites¹³. In this case it supports the coexistence of LDA and HDA phases for the amorphization of a hybrid system, so-called polyamorphic phases identical in composition but different in density and entropy. In contrast to ZIF-4, the structural integrity of ZIF-8 was maintained throughout the heating process (Fig. 2c,d).

Discussion

The DSC upscan curves of ZIF-4 (Fig. 3a, Supplementary Fig. 4) display endotherms (A) from release of framework templating *N,N*-dimethylformamide (DMF), which does not cause framework collapse. An exothermic feature follows (D–F), which indicates LDA T_g (Fig. 3a) (confirmed by SAXS experiments, Fig. 3c). Above this temperature, the resultant LDL converts to a HDL (Fig. 3a F–H)—corresponding to the HDA phase heated above T_g (Fig. 3d). This order–disorder transition is similar to polyamorphic transitions in inorganic zeolites^{4,13} and glass-forming liquids¹². Further heating results in recrystallization of HDL to ZIF-zni¹⁸. When the HDL is cooled to room temperature, after completion of the liquid–liquid transition (LLT), the HDA phase forms. This is confirmed by the occurrence of the glass transition at $T_g = 565$ K during reheating of the HDA, which is significantly lower than LDA $T_g = 589$ K (Fig. 3b), as found earlier in amorphizing inorganic zeolites. At the same time, the HDA–LDA LLT (H–C Fig. 3a) is not retraced at least with the DSC cooling rates currently available.

The various stages of amorphization are significantly heating rate dependent (Fig. 3b,c, Supplementary Fig. 4), from which the striking differences in fragility between HDL and LDL phases can be obtained (Fig. 4a). Remarkably, coexistence of LDA and HDA phases in the sample during amorphization is captured from double DSC scans (Fig. 3a,d), by progressively preheating to temperatures from desolvation, through collapse to the polyamorphic LDL–HDL transition (curves A–H in Fig. 3d). This can clearly be seen in curve D, where the endothermic response is followed by an exothermic one. The first relates to the

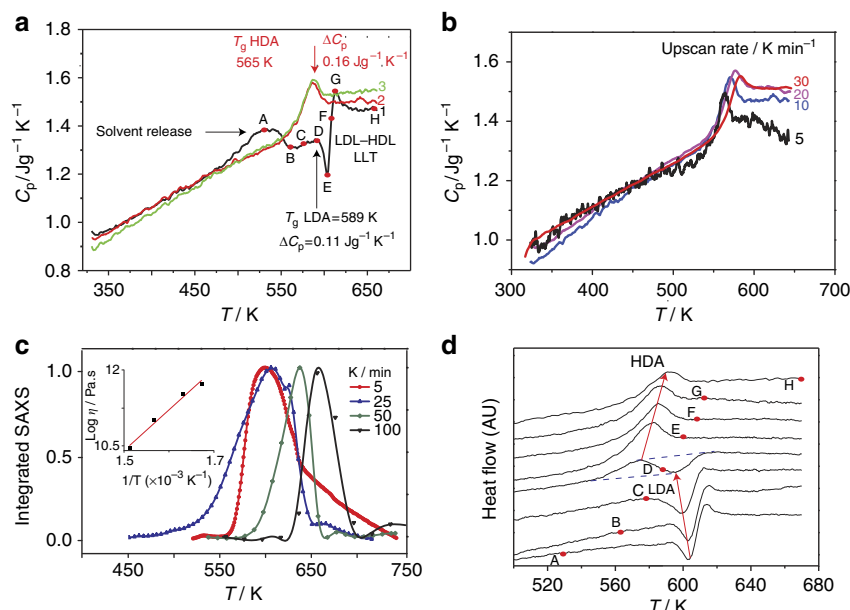


Figure 3 | Dynamics of ZIF-4 amorphization, polyamorphic glass transitions and coexistence (a) Sequence of DSC up-scans on ZIF-4 at 10 K min^{-1} starting with ZIF-4 (black), showing: solvent release (A), collapse to LDL phase (D–F), followed by the LLT to HDL (F–H). The jump in the isobaric heat capacity (C_p) through the LLT (E–G) is $0.33 \text{ J g}^{-1} \text{ K}^{-1}$. ΔC_p is the difference in C_p from glass to liquid at T_g , being 0.11 and $0.16 \text{ J g}^{-1} \text{ K}^{-1}$ for LDA and HDA phases, respectively. The endotherms in successive scans (2–red, 3–green) relate to HDA phase. (b) DSC second up-scans on the same samples at different rates right after cooling, yielding T_g and m for HDA. (c) The change in integrated SAXS ($\int_{0.04369}^{5.9253} I(q)_{\text{SAXS}} q^2 dq$), showing the increase of the peak temperature (T_{peak}) for different heating rates, giving T_g and m for the LDA phase. Inset: dependences of the Maxwell viscosity¹² $\eta = G_{\infty} \tau$, where G_{∞} and τ are the adiabatic shear modulus (2 GPa)⁴⁰ and structural relaxation time $\sim 1/\text{heating rate}$, respectively. (d) DSC up-scans preheated to temperatures A (529 K), B (563 K), C (578 K), D (588 K), E (601 K), F (608 K), G (613 K), H (673 K), cooled back to room temperature, and then reheated to 673 K—all at 10 K min^{-1} . Arrows indicate T_g HDA increasing and T_g LDA decreasing with increases in initial scan temperature. Temperature at 588 K reveals coexistence of LDA and HDA. With double scans (d), amorphization stages occur 20 K lower than for single scans (a).

T_g of the HDA phase already formed, with the second further collapse of ZIF-4 initiated at the T_g of LDA phase. The increase in the former follows the expected trend with increased pretreatment temperature, while the smaller but opposite trend in the latter suggests some increase in degrees of freedom as collapse advances. At coexistence (Fig. 3d) the 24 K difference between HDA and LDA glass transitions is reproduced. By comparison, single scanning (Fig. 3a), starting from ZIF-4 after solvent release, progresses consecutively through the respective transitions ZIF-4 to LDA (exothermic) and LDL–HDL (endothermic).

The large differences in viscosity of the glass-forming LDL and HDL phases can be quantified via Angell plots ($\log \eta$ versus T_g/T ; Fig. 4a), with respective fragilities of $m = 14$ and 41 resulting from use of structural relaxation times in SAXS (Fig. 3c) and DSC experiments (Supplementary Figs 5 and 6). Arrhenius ZIF-4 \rightarrow LDL collapse is hence what is expected for very strong liquids ($m = 14$), while HDL ($m = 41$) has intermediate fragility (Fig. 4a); this is in comparison to silica which is strong, the fragile anorthite and the very fragile triphenylethene. Given the melt fragility of silica ($m = 20$) (Fig. 4a), the LDL phase ($m = 14$) is referred to as a super-strong liquid^{8,12} and controls temperature induced collapse⁴.

ZIF-4 also collapses with pressure at room temperature between 0.35 GPa (P_1) and 0.98 GPa (P_2) (ref. 5), equivalent to thermal amorphization at ambient pressure between 603 K (T_1) and 638 K (T_2) (ref. 18). In accordance with prior work on zeolite instability, a T – P phase diagram similar in form to the two-liquid model of Rapoport^{4,24} is shown in Fig. 4b, constructed from P_1 , P_2 , T_1 and T_2 (refs 4,13). These are the pressure and temperature amorphization limits for the collapse of ZIF-4 and approximate to the spinodal limits for LDA and HDA phases. The negative

dT/dP slope and the increase in entropy through the LDA–HDA transition (Fig. 1b), as discussed earlier in the context of the Clapeyron relation and the amorphization of inorganic zeolites¹³, reaffirms the low and high densities of the ZIF-4 LDA and HDA phases. Furthermore, from the LDA–HDA excursion in C_p (E–G in Fig. 3a), the entropic rise between the two phases $\Delta S_{\text{LDA-HDA}}$ ($66 \text{ J mol}^{-1} \text{ K}^{-1}$) yields through $dT/dP = \Delta V_{\text{LDA-HDA}} / \Delta S_{\text{LDA-HDA}}$ a shrinkage of the molar volume of ZIF-4 (337 cm^3) $\Delta V_{\text{LDA-HDA}}$ of 10%. Following the line of enquiry in Fig. 4b, and extrapolating back the limits of the amorphization process, ZIF-4 LDA ($P_1 \rightarrow T_1$) and LDA–HDA ($P_2 \rightarrow T_2$), the LDA and HDA phases of ZIF-4 should become coexistent and identical at a critical point C at negative pressure, equivalent to what is observed in inorganic zeolites^{3,13}, and indeed similar to that predicted for amorphous silicon²⁵ and in yttria-alumina supercooled liquids²⁶. Critical points are associated with a sharp increase in density fluctuations²⁷, which, in Fig. 4b, will extend to ambient pressure, explaining the sharp peak in $I(q)_{\text{SAXS}}$ (Fig. 2a). The fact that the SAXS line shapes are closely Lorentzian is also consistent with the Ornstein–Zernike model for scattering close to critical points²⁸.

The chemical structures of ZIF-4, HDA and MQG phases probed by X-ray total scattering data are shown in Supplementary Fig. 11. All of the pair distribution functions (PDFs, Fig. 1c) below 6 \AA contain very similar sharp features, confirming the retention in HDA and MQG of the organic ligand and zinc tetrahedral coordination environments that characterize ZIF-4 (Fig. 1a). Given the similarity of MQG and HDA PDFs, along with the reconstructive transition from HDA to ZIF-zni¹⁸, we infer that some degree of Zn–N bond reconstruction occurs during amorphization and melting. Intriguingly, macroscopic

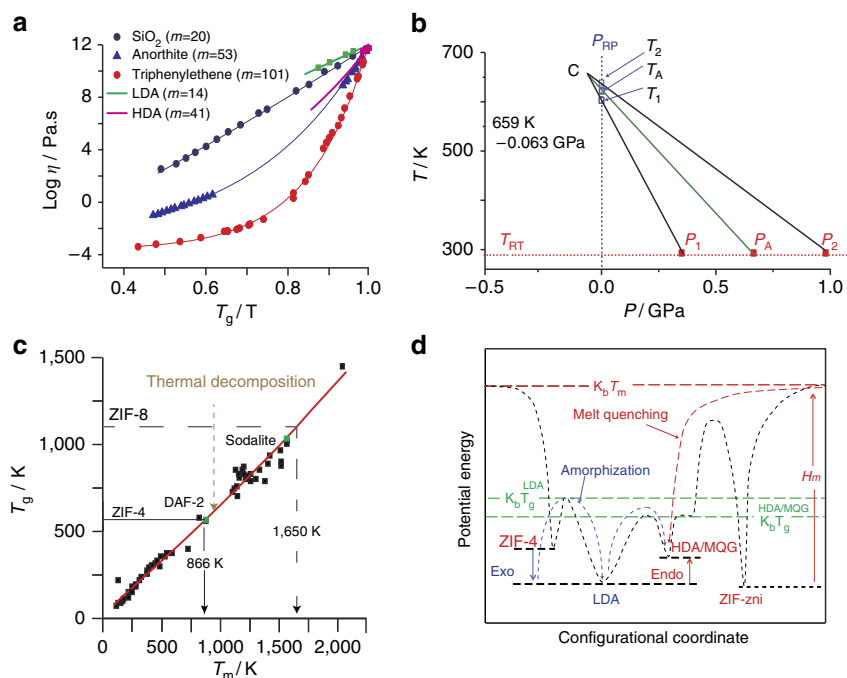


Figure 4 | Fragilities and critical point of ZIF-4 polyamorphs, projecting T_m from T_g and PEL schematic of ZIF-4 amorphization, melting and quenching routes. (a) Angell plot showing the fragility of LDL and HDL ZIF-4 (Fig. 3b,c), alongside other glass-forming liquids⁴¹ including the silica with <20 p.p.m. hydroxyl and <60 metallic impurities. Solid lines are fits to the measured viscosity-temperature relation of the model derived in previous literature^{16,42}. (b) T - P phase diagrams obtained from the limiting thermobaric amorphization parameters for ZIF-4 P_1 , P_2 , T_1 and T_2 , which extrapolate to a critical point C at negative pressure T_c (659 K) and P_c (-0.063 GPa). P_A and T_A refer to 50% amorphization points under pressure (RP)⁵ and temperature (RT)¹⁸, respectively. (c) 2/3's Law (T_g versus T_m) for different glass-forming systems^{29–31}, including ZIF-4 and ZIF-8 compared with DAF-2 and sodalite, respectively. The thermal degradation temperature separating the locations of the two amorphized ZIFs is shown. (d) Schematic of the PEL⁴³ for ZIF-4, informed from DSC experiments from Figs 1b and 3a. The adjacent LDA and HDA minima bear resemblance to the two states for water, different in density and topology, recently identified in modelling ST2 water³⁶.

flow of the melt into a non-porous glass (Brunauer-Emmett-Teller (BET) surface area of $<5 \text{ m}^2 \text{ g}^{-1}$) can be seen in scanning electron microscopy (SEM) and optical images (Fig. 1c, Supplementary Figs 7 and 8). The light brown color persists, even when oxygen is excluded from the reaction (Supplementary Fig. 9). At the same time ^1H NMR data recorded on digested samples confirm that imidazole ligands (Fig. 1a) remain largely intact (Supplementary Fig. 10).

The LDA glass transition temperature T_g (589 K) is extremely close to $2/3 T_m$ for ZIF = 4 (866 K), and therefore complies with the empirical law found for many glasses²⁹ (Fig. 4c). Interestingly, another MOF of $\text{Zn}(\text{Im})_2$ composition (ZIF-3), possesses the 'dft' zeolitic topology and undergoes identical amorphization and recrystallization to ZIF-4 (ref. 7). The inorganic cobalt phosphate framework DAF-2 (also adopting the 'dft' topology) is observed to melt at 873 K (ref. 22), indicating that frameworks with similar network topologies may exhibit similar melting behaviour, which may in turn be driven by collective THz modes⁶. In contrast, for Debye solids like dense minerals and metals, melting is activated by nearest neighbour r_{NN} vibrations when $\sqrt{(\Delta r_{\text{NN}}^2)/r_{\text{NN}}} \geq 0.1$ —Lindemann's Law²¹.

The 2/3's Law, already well-established for molecular and network glass formers^{30,31} and extended in Fig. 4c to include oxide glasses, enables T_m to be projected from the LDA T_g , where zeolites and MOFs collapse. In particular, despite ZIF-8 undergoing thermal decomposition before melting (Fig. 1b), a 'hypothetical' T_g of 1,100 K, can be calculated from the relationship $P_A \Delta V_A \approx 3RT_g$ (refs 4,7,23), where P_A is the amorphization collapse pressure and ΔV_A the collapsed volume. Using Fig. 4c this projects a 'virtual' T_m for ZIF-8 at 1,650 K. The temperature, which in practice is not achieved before

decomposition of the hybrid framework, lies close to T_m of the inorganic analogue (sodalite), which melts at 1,557 K. We postulate that, by this methodology, comparison of MOFs with their inorganic analogues should reveal candidates with achievable melting points and which could therefore form hybrid melt-quenched glasses like ZIF-4.

Fundamental to current understanding of the 2/3's Law^{30,31} is the kinetic fragility of the melt m and its association with the thermodynamic variables: heat of fusion, H_m and the jump of the isobaric heat capacity (C_p) from the glass at T_g to its liquid state above T_g , $\Delta C_p(T_g)$ viz, $\Delta C_p(T_g) = \frac{m H_m}{56 T_g}$. Recognizing the relationship between melt fragility and Poisson's ratio for the glass³², we have adapted this empirical relationship and recovered ΔC_p values (Fig. 3a) measured for LDL and HDL phases (Supplementary Methods, Supplementary Table 1). In addition, considering the various enthalpy changes occurring during amorphization, crystallization and melting (Fig. 1, Supplementary Fig. 3), a schematic PEL incorporating ZIF-4, LDA, HDA and MQG is plotted in Fig. 4d. It shows strong similarity with the behaviour of zeolite-A and its polyamorphs (Fig. 4d)¹³. It also illustrates the likely common location of HDA and MQG in configuration space, reached by amorphization and by melt quenching, respectively.

Finally, we consider the fact that, for these current experiments, cooling of the HDL ZIF phase does not result in formation of the LDL phase, encountered on heating (Fig. 3a). In terms of potential energy, the LDA ZIF phase shares similarities with the ultrastable glass obtained by molecule-by-molecule coating¹⁵. This also has a T_g greater than that of its 'normal' HDA glass counterpart. On reheating the ultrastable glass is transformed into the normal glass state, though does not convert back to the

ultrastable state within the cooling rates available using these techniques. This is also the same scenario observed in aged amber³³. By contrast, the LDA ZIF-4 phase reported here coexists with the HDA phase (Fig. 3d). When ZIF-4 has fully released its solvent, its structural arrangement becomes looser, but still remains ordered with an unchanged potential energy. With further heating the structure of solvent-free ZIF-4 relaxes towards a more stable state and lower enthalpy level, enthalpy being released with formation of the LDL phase (Figs 1b and 4d). This behaviour is also common to anhydrous zeolites^{4,13}, whose enthalpies all exceed those of conventional oxide glasses with the same composition³⁴, reflecting the metastable nature of zeolitic crystals. When the temperature subsequently rises above T_g , the LDL phase spontaneously transforms into the HDL ZIF phase. Beyond that, the HDL phase is finally turned to a more stable ZIF-zni crystal phase.

This apparent irreversibility for the LLT described here for the amorphization of ZIF-4 is in contrast to the reversibility reported in *ab initio* Molecular Dynamics volume versus pressure calculations on zeolites³⁵. With increasing pressure, zeolite-LDA followed by LDA-HDA first order transitions could also be retraced (albeit with some hysteresis) by reducing pressure, eventually including tension³⁵. Some evidence for phase transition reversibility was found experimentally during the initial zeolite collapsing process¹³. Elsewhere, experiments on yttria-alumina melts, where the LDL and HDL phases formed in levitated liquid drops, were observed to fluctuate back and forth at the LLT²⁶. Similar behaviour is confirmed both by the recent modelling of ST2 water³⁶ and experimental work on mannitol³⁷.

The apparent irreversibility of the LLT in ZIF-4 on current experimental timescales may lie kinetically in the inherent structural differences between hybrid and inorganic systems, in particular the comparative rigidity of the inter-tetrahedral bridging unit². Compared with oxide melts and zeolites, for example, the floppy bridging oxygen is replaced by the rigid imidazolate bridge in ZIF-4 (Fig. 1a). This will influence differences in conformational changes involved in the order-disorder LDA-HDA transition that determines the HDA topology, and may not be kinetically symmetric. So, while the entropy of the HDA phase lies comparatively close to that of the low-density state (Figs 1b and 3a), and both phases can coexist on heating (Fig. 3d), the difference is that for systems like water³⁶ and yttria-alumina melts²⁶, that are readily reversible, these appear to fluctuate freely between separate free energy basins facilitated by pivotal inter-polyhedral bridges. In particular, where the LDL to HDL transition in ZIF-4 is thermodynamically controlled, the reverse process appears to be kinetically controlled, the dynamics being out of range using current experimental methods.

Comparisons between amorphization and melting conditions of MOFs and inorganics may provide further routes to more functional 'perfect' glasses, HDA and MQG phases. Furthermore, the *in situ* hybrid liquid formation discovered here opens up possibilities for liquid casting and shaping MOFs into a variety of different solid forms, promising to be an extremely exciting step forward in producing chemically functionalizable hybrid glass materials.

Methods

Synthesis. 1.2 g of $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ and 0.9 g of imidazole were dissolved in 90 ml of DMF and transferred into a 100 ml screw jar. The jar was tightly sealed and heated to 100 °C for 72 h in an oven. After cooling to room temperature colourless block-shaped crystals were filtered off and first washed three times with ~30 ml pure DMF and then three times with ~30 ml CH_2Cl_2 . The HDA and quenched ZIF-zni used for the PDF experiments were formed by heating ZIF-4 to 573 and 865 K under an argon atmosphere using a ramp rate of 5 K min⁻¹.

To investigate the effect of oxygen on the process, a 1 mm diameter quartz tube was loaded with a sample of crystalline ZIF-4, and sealed under vacuum. The capillary was then heated in a tube furnace under an argon flow, at a rate of 5 °C min⁻¹, to 865 K. The final melt-quenched glass was not observed to differ from that attained in other experiments.

For SAXS and WAXS measurements, the crystals were gently stirred in 100 ml fresh CH_2Cl_2 overnight. Afterwards the solid material was filtered off, washed again three times with ~30 ml fresh CH_2Cl_2 and dried in vacuo at 130 °C, using a vacuum oven to yield activated guest-free ZIF-4.

ZIF-8 was purchased from Sigma Aldrich and evacuated by heating at 100 °C for 3 h.

Measurements. Room temperature X-ray total scattering data were collected at the I15 beamline at Diamond Light Source, UK, at a wavelength of $\lambda = 0.1722 \text{ \AA}$. Finely powdered samples of ZIF-4, HDA and MQG samples were carefully loaded into 1.0 mm diameter fused silica capillaries, and data from an empty instrument and capillary were also collected for background subtraction. Data were collected between $\sim 0.5 < Q < \sim 22 \text{ \AA}^{-1}$. Corrections for background, multiple scattering, container scattering, Compton scattering and absorption were applied. The normalized reciprocal space data (Supplementary Fig S1) were then converted to the PDFs using Fourier transform.

Temperature dependent *in situ* SAXS and WAXS measurements were performed on Beamline I22 at the Diamond Light Source synchrotron in the Rutherford Appleton Laboratory (Didcot, Oxfordshire, UK). Detector calibrations were carried out using NBS (National Bureau of Standards) silicon and silver behenate standards on the HOTWAXS 1D WAXS, and RAPID 2D SAXS, detectors respectively^{38,39}.

Normalization for beam intensity and sample thickness and/or density variation was carried out using a diode embedded in the beamstop. Scattering data were recorded at a wavelength of 1 Å for angular range of up to 1° for SAXS and over a 2θ range of 5°–40° for WAXS. Powdered samples of ZIF-4 and ZIF-8 were loaded in glass capillaries and inserted horizontally through the Linkam furnace, which was positioned across the synchrotron radiation source. The Linkam furnace used was calibrated, finding the relationship $T_{\text{true}} = 0.95 T (\text{set point, C}) + 65$. Raw data for temperature scanned SAXS ZIF-4 and ZIF-8 are shown in Supplementary Fig. 2. SAXS profiles were fitted to Lorentzian line shapes, $I(q) = \frac{I(0)}{1 + q^2\xi^2}$, with the correlation length ξ increasing from ~220 to ~840 Å from the edge to the peak (Fig. 2a). Additional measurements accompanied by a detailed analysis will be given in a separate publication.

The apparent isobaric heat capacity (C_p) of each sample was measured using a Netzsch STA 449C DSC. The samples were placed in a platinum crucible situated on a sample holder of the DSC at room temperature and subjected to varying numbers of up- and down-scans, depending on the purpose of the measurements. After natural cooling to room temperature, the subsequent up-scans were performed using the same procedure as for the first.

Powder X-ray diffraction measurements on evacuated, guest-free ZIF-4 were recorded on a well ground sample with a Bruker D8 Advance powder diffractometer using $\text{CuK}\alpha$ radiation ($\lambda = 1.5418 \text{ \AA}$) and a LynxEye position sensitive detector in Bragg-Brentano ($\theta - \theta$) geometry at room temperature. Pawley fit shown in Supplementary Fig. 11.

Microanalysis was performed at the Department of Chemistry, University of Cambridge as a technical service.

ZIF-4 evacuated. Calculated (based on $\text{Zn}(\text{C}_3\text{H}_3\text{N}_2)_2$ composition): C 36.18%, H 3.02%, N 28.14%. Found: C 36.22%, H 2.98%, N 28.09%

MOF glass. Calculated (based on $\text{Zn}(\text{C}_3\text{H}_3\text{N}_2)_2$ composition): C 36.18%, H 3.02%, N 28.14%. Found: C 35.64%, H 2.90%, N 26.46%

SEM images (Supplementary Fig. 7) were taken with an FEI Nova NanSEM (field emission gun). Specimens for SEM analysis were prepared by dispersing fragments of the ZIF-4 melt-quenched glass on conductive carbon tabs for topographic contrast imaging. Optical images of ZIF-4, ZIF-zni and recovered melt-quenched glass are shown in Supplementary Fig. 8.

Liquid phase ^1H NMR spectra (Supplementary Fig. 10) of digested samples ($\text{DCl}/\text{D}_2\text{O}/\text{DMSO}-d_6$) of evacuated ZIF-4 and the ZIF-4 glass (~5–10 mg) were recorded on a Bruker Avance DPX-250 spectrometer at 293 K in a mixture of DCl (35%)/ D_2O (0.1 ml) and $\text{DMSO}-d_6$ (0.5 ml). Chemical shifts are given relative to tetramethylsilane and were referenced to the residual protio-solvent signals of $\text{DMSO}-d_6$. The spectra were processed with the MestreNova Suite.

Nitrogen adsorption Brunauer-Emmett-Teller (BET) measurements were carried out at 77 K using a Micromeritics ASAP 2020 instrument.

References

1. Furukawa, H., Cordova, K. E., O'Keeffe, M. & Yaghi, O. M. The chemistry and applications of metal-organic frameworks. *Science* **341**, 974–986 (2013).
2. Rogez, G., Viart, N. & Drillon, M. Multiferroic materials: the attractive approach of metal-organic frameworks (MOFs). *Angew. Chem. Int. Ed.* **49**, 1921–1923 (2010).
3. Bennett, T. D. & Cheetham, A. K. Amorphous metal-organic frameworks. *Acc. Chem. Res.* **47**, 1555–1562 (2014).
4. Greaves, G. N. *et al.* The rheology of collapsing zeolites amorphized by temperature and pressure. *Nat. Mater.* **2**, 622–629 (2003).

5. Bennett, T. D. *et al.* Reversible pressure-induced amorphization of a zeolitic imidazolate framework (ZIF-4). *Chem. Commun.* **47**, 7983–7985 (2011).
6. Greaves, G. N., Meneau, F., Majerus, O., Jones, D. G. & Taylor, J. Identifying vibrations that destabilize crystals and characterize the glassy state. *Science* **308**, 1299–1302 (2005).
7. Bennett, T. D. *et al.* Thermal amorphization of zeolitic imidazolate frameworks. *Angew. Chem. Int. Ed.* **50**, 3067–3071 (2011).
8. Angell, C. A., Moynihan, C. T. & Hemmati, M. 'Strong' and 'superstrong' liquids, and an approach to the perfect glass state via phase transition. *J. Non-Cryst. Solids* **274**, 319–331 (2000).
9. Mishima, O., Calvert, L. D. & Whalley, E. An apparently first-order transition between two amorphous phases of ice induced by pressure. *Nature* **314**, 76–78 (1985).
10. Deb, S. K., Wilding, M., Somayazulu, M. & McMillan, P. F. Pressure-induced amorphization and an amorphous-amorphous transition in densified porous silicon. *Nature* **414**, 528–530 (2001).
11. Ding, S. P. *et al.* Vitrification of trehalose by water loss from its crystalline dihydrate. *J. Therm. Anal.* **47**, 1391–1405 (1996).
12. Greaves, G. N. & Sen, S. Inorganic glasses, glass-forming liquids and amorphizing solids. *Adv. Phys.* **56**, 1–166 (2007).
13. Greaves, G. N. *et al.* Zeolite collapse and polyamorphism. *J. Phys.-Condens. Mat.* **19**, 415102 (2007).
14. Haines, J. *et al.* Topologically ordered amorphous silica obtained from the collapsed siliceous zeolite, silicalite-1-F: a step toward "Perfect" glasses. *J. Am. Chem. Soc.* **131**, 12333–12338 (2009).
15. Singh, S., Ediger, M. D. & de Pablo, J. J. Ultrastable glasses from in silico vapour deposition. *Nat. Mater.* **12**, 139–144 (2013).
16. Angell, C. A. Formation of glasses from liquids and biopolymers. *Science* **267**, 1924–1935 (1995).
17. Amann-Winkel, K. *et al.* Water's second glass transition. *Proc. Natl Acad. Sci. USA* **110**, 17720–17725 (2013).
18. Bennett, T. D. *et al.* Structure and properties of an amorphous metal-organic framework. *Phys. Rev. Lett.* **104**, 115503 (2010).
19. Angell, A. in *The International Symposium on Non Oxide and New Optical Glasses* (Saint Malo, 2012).
20. Park, K. S. *et al.* Exceptional chemical and thermal stability of zeolitic imidazolate frameworks. *Proc. Natl Acad. Sci. USA* **103**, 10186–10191 (2006).
21. Poirer, J. P. *Introduction to the Physics of the Earth's Interior* (Cambridge University Press, 2000).
22. Chen, J. S., Jones, R. H., Natarajan, S., Hursthouse, M. B. & Thomas, J. M. A novel open-framework cobalt phosphate-containing a tetrahedrally coordinated Cobalt(II) center - $\text{CoPO}_4 \cdot 0.5 \text{C}_2\text{H}_{10}\text{N}_2$. *Angew. Chem. Int. Ed.* **33**, 639–640 (1994).
23. Chapman, K. W., Halder, G. J. & Chupas, P. J. Pressure-induced amorphization and porosity modification in a metal-organic framework. *J. Am. Chem. Soc.* **131**, 17546–17547 (2009).
24. Rapoport, E. Model for melting curve maxima at high pressure. *J. Chem. Phys.* **46**, 2891–2895 (1967).
25. Aptekar, L. I. Phase-transition in noncrystalline germanium and silicon. *Sov. Phys. Dokl.* **249**, 1099–1103 (1979).
26. Greaves, G. N. *et al.* Detection of first-order liquid/liquid phase transitions in yttrium oxide-aluminum oxide melts. *Science* **322**, 566–570 (2008).
27. Stanley, H. E. *Introduction to Phase Transitions and Critical Phenomena* (Oxford University Press, 1971).
28. Ornstein, L. S. & Zernike, F. Accidental deviations of density and opalescence at the critical point of a single substance. *Proc. Acad. Sci. Amst.* **17**, 793–806 (1914).
29. Kauzmann, W. The nature of the glassy state and the behaviour of liquids at low temperatures. *Chem. Rev.* **43**, 219–256 (1948).
30. Wang, L. M., Angell, C. A. & Richert, R. Fragility and thermodynamics in nonpolymeric glass-forming liquids. *J. Chem. Phys.* **125**, 074505 (2006).
31. Stevenson, J. D. & Wolynes, P. G. Thermodynamic-kinetic correlations in supercooled liquids: a critical survey of experimental data and predictions of the random first-order transition theory of glasses. *J. Phys. Chem. B* **109**, 15093–15097 (2005).
32. Novikov, V. N. & Sokolov, A. P. Poisson's ratio and the fragility of glass-forming liquids. *Nature* **431**, 961–963 (2004).
33. Perez-Castaneda, T., Rodriguez-Tinoco, C., Rodriguez-Viejo, J. & Ramos, M. A. Suppression of tunneling two-level systems in ultrastable glasses of indomethacin. *Proc. Natl Acad. Sci. USA* **111**, 11275–11280 (2014).
34. Navrotsky, A. & Tian, Z. R. Systematics in the enthalpies of formation of anhydrous aluminosilicate zeolites, glasses, and dense phases. *Chem.-Eur. J.* **7**, 769–774 (2001).
35. Peral, I. & Iniguez, J. Amorphization induced by pressure: results for zeolites and general implications. *Phys. Rev. Lett.* **97**, 225502 (2006).
36. Palmer, J. C. *et al.* Metastable liquid-liquid transition in a molecular model of water. *Nature* **510**, 385–388 (2014).
37. Zhu, M., Wang, J.-Q., Perepezko, J. H. & Yu, L. Possible existence of two amorphous phases of d-mannitol related by a first-order transition. *J. Chem. Phys.* **142**, 244504 (2015).
38. Bateman, J. E. *et al.* The HOTWAXS detector. *Nucl. Instrum. Methods A* **580**, 1526 (2007).
39. Hall, C. J., Lewis, R. A., Berry, A., Helsby, W. I. & Parker, B. T. The RAPID detector system - first user data. *Nucl. Instrum. Methods A* **454**, 2000 (2000).
40. Tan, J. C., Civalleri, B., Erba, A. & Albanese, E. Quantum mechanical predictions to elucidate the anisotropic elastic properties of zeolitic imidazolate frameworks: ZIF-4 vs ZIF-zni. *CrystEngComm* **17**, 375–382 (2015).
41. Debenedetti, P. G. & Stillinger, F. H. Supercooled liquids and the glass transition. *Nature* **410**, 259–267 (2001).
42. Mauro, J. C., Yue, Y. Z., Ellison, A. J., Gupta, P. K. & Allan, D. C. Viscosity of glass-forming liquids. *Proc. Natl Acad. Sci. USA* **106**, 19780–19784 (2009).
43. Angell, A. Thermodynamics - Liquid landscape. *Nature* **393**, 521–524 (1998).

Acknowledgements

The authors would like to thank Trinity Hall (T.D.B.); HRH Sheikh Saud Bin Saqr Al Qasimi (T.D.B. and A.K.C.); Wuhan University of Science and Technology (Y.Z.Y. and G.N.G.), ERC grant number 259619 PHOTO EM (C.D.); the Alexander von Humboldt Foundation (S.H.); The World Premier International Research Center Initiative on Materials Nanoarchitectonics (WPI-MANA) from MEXT, Japan (H.H.M.Y.). We acknowledge the provision of synchrotron access to Beamline I15 (exp. EE9691) and I22 (exp. SM5692) at the Diamond Light Source, Rutherford Appleton Laboratory UK. Dr Philip Chater (Diamond Light Source) and Andrew Cairns (Oxford) are thanked for assistance during the beamtime on I15, as is Professor David Keen (ISIS) for valuable assistance with the data analysis. Professor Seth Cohen and Xiao Yu (University California San Diego) are acknowledged for BET measurements.

Author contributions

G.N.G. with A.K.C. facilitating MOF materials and characterization. T.D.B., Y.Z.Y. and G.N.G. wrote the manuscript, J.C.T. with A.K.C. involved in refinements. J.C.T., H.H.M.Y., N.T. and G.N.G. carried out the *in situ* SAXS and WAXS experiments at the Diamond Light Source; J.C.T. analysed the synchrotron data and established the melt fragilities guided by G.N.G. Y.Z.Y. performed the DSC measurements and data analysis. S.H. synthesized and characterized ZIF-4 crystals for DSC and melting experiments, and performed NMR experiments. T.D.B. carried out PXRD, SDT and FTIR measurements on ZIF-4, and synthesized and performed elemental analysis and optical microscopy on recovered MQG. C.D. performed SEM imaging of MOF glass. W.C. and Z.Z. contributed to visualization and graphics, including modelling amorphized ZIF-8. T.D.B. and S.H. involved in the preparation of powder samples used in this study. T.D.B. led the PDF measurements, which were carried out with E.B.

Additional information

Supplementary Information accompanies this paper at <http://www.nature.com/naturecommunications>

Competing financial interests: The authors declare no competing financial interests.

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How to cite this article: Bennett, T. D. *et al.* Hybrid glasses from strong and fragile metal-organic framework liquids. *Nat. Commun.* 6:8079 doi: 10.1038/ncomms9079 (2012).



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